ABSTRACT

There is provided a T-type calcium channel blocker that is a compound of formula (1), a pharmaceutically acceptable salt thereof or a solvate thereof:

$$Z \xrightarrow{Ar^1} CO_2R^3$$

$$R^b \qquad (1)$$

wherein

Ar¹ is phenyl group, pyridyl group, furyl group or 2,1,3-benzoxadiazol-4-yl group; nitrogen-containing hetero ring moiety is 1,4-dihydropyridine ring or pyridine ring; Z is a group of formula (2)

or CO₂R²;

R^a and R^b are independently of each other C_{1-a}alkyl group, ANR^aR^a, CH₂OANR^aR^a, or the like;

in case where the nitrogen-containing hetero ring molety is 1,4-dihydropyridine ring, R¹ is C₁₋₈alkyl group, ANR⁸R⁹, AN(CH₂CH₂)₂NR⁸, AN(CH₂CH₂)₂O, AOR⁸ or benzyl group;

R³ is hydrogen atom, C₁₋₂₀alkyl group, ANR®R9, a group of formula

$$-A-N$$
 $N-R^8$, $-A-N$ $N-R^6$, $N-R^6$

or the like.